1. (Original) A method of determining the presence of a nuclear localization signal in a protein of interest, the method comprising:

selecting a host cell for use in the method, wherein the host cell contains a nucleus having nucleic acid encoding a reporter gene therein and wherein the host cell has a first level of expression of the reporter gene;

identifying a DNA binding domain and an activation domain for the reporter gene;

constructing a chimeric nucleic acid encoding a fusion protein comprising the DNA binding domain, the activation domain, and a protein of interest, wherein elements of the fusion protein other than the protein of interest have no nuclear localization signals;

introducing the chimeric nucleic acid into the host cell; and

determining a second level of expression of the reporter gene to determine the presence of a nuclear localization signal in the protein of interest.

- 2. (Original) The method of claim 1 wherein the host cell is a eukaryotic cell.
- 3. (Original) The method of claim 1 wherein the host cell is a yeast cell.
- 4. (Original) The method of claim 1 wherein the reporter gene is a lacZ gene.
- 5. (Original) The method of claim 1 wherein the reporter gene is a selection marker gene.
- 6. (Original) The method of claim 5 wherein the selection marker gene is a HIS3 gene.

- 7. (Previously Amended) The method of claim 4 or 6 wherein the DNA binding domain is from a LexA protein.
- 8. (Original) The method of claim 4 or 6 wherein the activation domain is a GAL4 activation domain.
- 9. (Original) The method of claim 1 wherein the chimeric nucleic acid further comprises nucleic acid encoding a promoter to control expression of the fusion protein.
- 10. (Original) The method of claim 9 wherein the promoter is an ADH1 promoter.
- 11. (Currently Amended) A recombinant host cell comprising:
- a nucleus having nucleic acid encoding a reporter gene therein; and
- a chimeric nucleic acid encoding a fusion protein, the fusion protein comprising a DNA binding domain for the reporter gene, an activation domain for the reporter gene, and a protein of interest, wherein elements of the fusion protein other than the protein of interest have no nuclear localization signals and wherein the DNA binding domain is from a LexA protein.
- 12. (Original) The recombinant host cell of claim 11 wherein the host cell is a eukaryotic cell.
- 13. (Original) The recombinant host cell of claim 11 wherein the host cell is a yeast cell.
- 14. (Original) The recombinant host cell of claim 11 wherein the reporter gene is a lacZ gene.
- 15. (Original) The recombinant host cell of claim 11 wherein the reporter gene is a selection marker gene.

- 16. (Original) The recombinant host cell of claim 15 wherein the selection marker gene is a HIS3 gene.
 - 17. (Canceled)
- 18. (Original) The recombinant host cell of claim 14 or 16 wherein the activation domain is a GAL4 activation domain.
- 19. (Original) The recombinant host cell of claim 11 wherein the chimeric nucleic acid further comprises nucleic acid encoding a promoter to control expression of the fusion protein.
- 20. (Original) The recombinant host cell of claim 19 wherein the promoter is an ADH1 promoter.
- 21. (Original) A chimeric nucleic acid encoding a fusion protein, the fusion protein comprising a DNA binding domain for a reporter gene, an activation domain for the reporter gene, and a protein of interest, wherein elements of the fusion protein other than the protein of interest have no nuclear localization signals and wherein the DNA binding domain is from a LexA protein.
- 22. (Original) The chimeric nucleic acid of claim 21 wherein the reporter gene is a lacZ gene.
- 23. (Original) The chimeric nucleic acid of claim 21 wherein the reporter gene is a selection marker gene.
- 24. (Original) The chimeric nucleic acid of claim 23 wherein the selection marker gene is a HIS3 gene.
 - 25. (Canceled)

- 26. (Original) The chimeric nucleic acid of claim 22 or 24 wherein the activation domain is a GAL4 activation domain.
- 27. (Original) The chimeric nucleic acid of claim 21 further comprising nucleic acid encoding a promoter to control expression of the fusion protein.
- 28. (Original) The chimeric nucleic acid of claim 27 wherein the promoter is an ADH1 promoter.
- 29. (Original) A vector comprising the chimeric nucleic acid of claim 21.
 - 30. (Original) A kit comprising the vector of claim 29.
- 31. (Original) The kit of claim 30 further comprising host cells which contain a nucleus having nucleic acid encoding the reporter gene therein.
- 32. (Original) The kit of claim 31 further comprising a control vector.
- 33. (Original) A nucleic acid molecule encoding a modified LexA protein, wherein the modified LexA protein has no nuclear localization signal.
- 34. (Original) The nucleic acid molecule of claim 33 wherein the nucleic acid molecule has a nucleotide sequence as shown in SEQ ID NO:1.
- 35. (Original) The nucleic acid molecule of claim 33 wherein the nucleic acid molecule encodes an amino acid sequence as shown in SEQ ID NO:2.
- 36. (Original) A modified LexA protein, wherein the modified LexA protein has no nuclear localization signal.

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37. (Original) The modified LexA protein of claim 36 wherein the protein has an amino acid sequence as shown in SEQ ID NO:2.

38.-78. (Canceled)